



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application of:
Mary Ellen Rybak, et al.

Serial No.: 09/904,263

Filed: July 12, 2001

For: MELANOMA THERAPY

Examiner: Anne L. Holleran

Group Art Unit: 1642

Atty. Docket No.: OC01000KQ

SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT

Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Sir:

(a) Pursuant to Rule 56, it is requested that the documents listed on the accompanying PTO-1449 Form be considered and made of record in the above-identified patent application. Copy(ies) of these references ☒ are attached ☐ were filed in related application U.S. Serial No(s) _____ filed _____, respectively.

(b) ☒ No fee is believed due because:

- ☐ This Information Disclosure Statement is being filed within three (3) months of the filing date of the application.
- ☐ This Information Disclosure Statement is being concurrently filed with the above-identified application.
- ☒ This Information Disclosure Statement is being concurrently filed with a Request for Continued Examination (RCE).
- ☐ This Information Disclosure Statement is being filed prior to the mailing of a first Office Action on the merits.

(c) ☐ This Information Disclosure Statement is being filed before the mailing date of any final action, notice of allowance or an action that otherwise closes prosecution; and

- ☐ Each item of information contained in this Information Disclosure Statement was first cited in a communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of this Information Disclosure Statement; or
- ☐ No item of information contained in this Information Disclosure Statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the undersigned after making reasonable inquiry, no item of information contained in this Information Disclosure Statement was known to any individual designated in § 1.56(c) more than three months prior to the filing of this Information Disclosure Statement; or

☐ The Commissioner is hereby authorized to charge the requisite fee listed on the attached Fee Transmittal Sheet.

(d) ☐ This Information Disclosure Statement is being filed on or before the payment of the issue fee; and

- ☐ Each item of information contained in this Information Disclosure Statement was first cited in a communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of this Information Disclosure Statement; or
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- ☐ The Commissioner is hereby authorized to charge the requisite fee listed on the attached Fee Transmittal Sheet.
- ☒ The Commissioner is hereby authorized to charge any additional fees which may be required for this Information Disclosure Statement, or credit any overpayment to Deposit Account No. **19-0365**, Patent Case No. **OC01000KQ**. A DUPLICATE COPY OF THIS SHEET IS ATTACHED.

Respectfully submitted,

SCHERING-PLOUGH CORPORATION

Dated: March 22, 2004

By:

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Sheet 1 of 1

FORM PTO-1449		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTY. DOCKET NO.: OC01000KQ		APPLICATION NO.: 09/904,263	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use several sheets if necessary)				APPLICANT: Mary Ellen Rybak, et al.			
				FILING DATE: 07/12/2001		GROUP: 1642	
U.S. PATENT DOCUMENTS							
*EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUB- CLASS	FILING DATE IF APPROPRIATE
FOREIGN PATENT DOCUMENTS							
		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB- CLASS	TRANSLATION YES NO
OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)							
	AA	Bruno, René, <i>et al.</i> , "Population Pharmacokinetics/Pharmacodynamics of Docetaxel in Phase II Studies in Patients with Cancer," <i>Journal of Clinical Oncology</i> 16 (1):187-96 (1998)					
	AB	Calvert, A.H., <i>et al.</i> , "Carboplatin Dosage: Prospective Evaluation of a Simple Formula Based on Renal Function," <i>Journal of Clinical Oncology</i> 7 (11):1748-56 (1989)					
	AC	Eisenhauer, Elizabeth A., <i>et al.</i> , "The Taxoids, Comparative Clinical Pharmacology and Therapeutic Potential," <i>Drugs</i> 55 (1):5-30 (1998)					
	AD	Eksborg, Steffan, <i>et al.</i> , "Plasma pharmacokinetics of idarubicin and its 13-dihydro metabolite-a comparison of bolus versus 2 h infusion during a 3 day course," <i>Anti-Cancer Drugs</i> 8 :42-7 (1997)					
	AE	Forastiere, Arlene A., <i>et al.</i> , "Pharmacokinetic and Toxicity Evaluation of Five-Day Continuous Infusion versus Intermittent Bolus <i>cis</i> -Diamminedichloroplatinum(II) in Head and Neck Cancer Patients," <i>Cancer Research</i> 48 :3869-3874 (1988)					
	AF	Gandhi, Varsha, <i>et al.</i> , "Compound GW506U78 in Refractory Hematologic Malignancies: Relationship Between Cellular Pharmacokinetics and Clinical Response," <i>Journal of Clinical Oncology</i> 16 (11):3607-15 (1998)					
	AG	Rodman, John H., <i>et al.</i> , "Clinical Pharmacodynamics of Continuous Infusion Teniposide: Systemic Exposure as a Determinant of Response in a Phase I Trial," <i>Journal of Clinical Oncology</i> 5 (7):1007-1014 (1987)					
	AH	Shalinsky, David R., <i>et al.</i> , "Antitumor efficacy of AG3340 associated with maintenance of minimum effective plasma concentrations and not total daily dose, exposure or peak plasma concentrations," <i>Investigational New Drugs</i> 16 :303-13 (1999)					
	AI	Takitani, Kimitaka, <i>et al.</i> , "4-Oxo Retinoic Acid for Refractory Acute Promyelocytic Leukemia in Children with All-Trans Retinoic Acid Therapy," <i>J. Nutr. Sci. Vitaminol.</i> 41 :493-98 (1995)					
EXAMINER				DATE CONSIDERED			
*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.							